

Bovilis BVD Suspension for injection for cattle

Authorised

- Bovine viral diarrhoea virus 1, strain C-86, Inactivated

Product identification

Medicine name:

Bovilis BVD Suspension for injection for cattle

Active substance:

Bovine viral diarrhoea virus 1, strain C-86, Inactivated

Target species:

Cattle

Cattle (for meat production)

Cattle (dairy cow)

Cattle (calf)

Cattle (heifer)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Bovine viral diarrhoea virus 1, strain C-86, Inactivated

50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Cattle

- Milk. no withdrawal period zero days

- Meat and offal. no withdrawal period zero ays

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Cattle (for meat production)

- Meat and offal. no withdrawal period zero days

-

Cattle (dairy cow)

- Milk. no withdrawal period zero days

- Meat and offal. no withdrawal period zero days

-

Cattle (calf)

- Meat and offal. no withdrawal period zero days

-

Cattle (heifer)

- Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

(ID7) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 100 millilitre(s)

(ID6) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)

(ID1) 10 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 millilitre(s)

(ID20) 2 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 2 millilitre(s)

(ID30) 2 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 2 millilitre(s)

(ID3) 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 20 millilitre(s)

(ID9) 250 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 250 millilitre(s)

(ID4) 20 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)

(ID2) 10 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 10 millilitre(s)

(ID5) 50 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 50 millilitre(s)

(ID8) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)

(ID10) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

13/09/1999

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

13/09/1999

Reference member state:

Netherlands

Procedure number:

NL/V/0433/001

Concerned member states:

Austria Belgium France Germany Greece Ireland Italy Luxembourg Poland
Portugal Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 28/01/2022

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Package Leaflet

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Labelling

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